12 510(K) SUMMARY OF SAFETY AND EFFECTIVENESS

K001841

Submission in accordance with the requirements of 21 CFR Part 807.87(h)

1. Submitter

Applicant:	**TAGARNO A/S** Hattingvej 5 DK-8700 Horsens Denmark
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Manufacturer:	**TAGARNO A/S** Hattingvej 5 DK-8700 Horsens Denmark
Owner/operator ID number:	9040414
Prepared:	16. June 2000

2. Device Name

Common name: 35 / CD CINECONVERTER
Classification name: Digitizer, Image Processing
Regulation Number: 892.2030, LMA 414

3. Predicate Device

OSCAR, Arri, K993283 (=substantial equivalence)

4) Description of the device:

The **TAGARNO** 35/CD CINECONVERTER system consist of three major components, the **TAGARNO** 35/CD cine film projector, a PC equipped with a frame grabber and a CD writer and the **TAGARNO** View software package.

5) Intended use

The **TAGARNO** 35/CD CINECONVERTER system will be used to digitize X-ray angio 35 mm. cine films and export those digitized cine films in DICOM V3 format.

6) Substantial equivalence Information

TAGARNO 35/CD CINECONVERTER is substantially equivalent to the Predicate Devices of ARRI, Oscar K993283.

Conclusion respecting safety and effectiveness:
It is the opinion of **TAGARNO** that the Cineconverter is safe and potential hazards are controlled by a risk management plan for the development process, including hazard analysis, verification and tests (see Appendix 2). The software package itself will not have any adverse effects on health.
In **TAGARNO**'s opinion the level of concern for this device to view images is 'minor' and that the use of this device supports the clinician in their decision process; nor does the use of the device result in any new potential hazards.

Horsens, 16. June 2000

Max Jens Jensen
Plant manager
TAGARNO A/S

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

NOV 1 5 2000

Mr. Max Jens Jensen
Plant Manager
TAGARNO A/S
Hattingvej 5
DK-8700 Horsens
DENMARK

Re: K001841
TAGARNO 35/CD CINECONVERTER System
Dated: September 19, 2000
Received: September 21, 2000
Regulatory Class: II
21 CFR §892.2030/Procode: 90 LMA

Dear Mr. Jensen:

We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the Current Good Manufacturing Practice requirements, as set forth in the Quality System Regulation (QS) for Medical Devices: General regulation (21 CFR Part 820) and that, through periodic QS inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your device in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

This letter will allow you to begin marketing your device as described in your 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4639. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its internet address "http://www.fda.gov/cdrh/dsma/dsmamain.html".

Sincerely yours,

Daniel G. Schultz, M.D.
Captain, USPHS
Acting Director, Division of Reproductive,
Abdominal, and Radiological Devices
Office of Device Evaluation
Center for Devices and Radiological Health

Enclosure (s)

510(k) Number (if known): K001841

Device Name: 35/CD CINECONVERTER

Indications For Use:

The **TAGARNO** 35/CD CINECONVERTER system will be used to digitize X-ray angio 35 mm. cine films and export those digitized cine films in DICOM V3 format.

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

(Division Sign-Off)
Division of Reproductive, Abdominal, ENT,
and Radiological Devices
510(k) Number K001841

Prescription Use ✓ OR Over-The-Counter Use ___
(Per 21 CFR 801.109)

(Optional Format 1-2-96)